Amendments to the claims:

1. (original) A pharmaceutical composition for treatment of pain, which comprises in combination a benzothiadiazole derivative of formula I

wherein each R1, R2 and R3 independently, is hydrogen, halogen, C_1 - C_7 alkyl, C_1 - C_7 alkoxy, nitro, cyano, hydroxy or C_1 - C_7 alkylthio; and a COX-2 inhibitor for simultaneous, sequential or separate use.

- 2. (canceled)
- 3. (canceled)
- 4. (currently amended) A method of treating a patient suffering from pain comprising administering to the patient an effective amount of a benzothiadiazole derivative of formula I as defined above.

wherein each R1, R2 and R3 independently, is hydrogen, halogen, C_1 - C_7 alkyl, C_1 - C_7 alkoxy, nitro, cyano, hydroxy or C_1 - C_7 alkylthio, and an effective amount of a COX-2 inhibitor.

5. (currently amended) A package comprising a benzothiadiazole derivative of formula I as defined in claim 1,

wherein each R1, R2 and R3 independently, is hydrogen, halogen, C₁-C₇ alkyl, C₁-C₇ alkoxy, nitro, cyano, hydroxy or C₁-C₇ alkylthio, for treatment of pain according to a method wherein said benzothiadiazole derivative of formula I is used together with instructions for use in combination with a COX-2 inhibitor for treatment of pain, or a package comprising a COX-2 inhibitor for treatment of pain according to a method wherein said COX-2 inhibitor is used together with instructions for use in combination with a benzothiadiazole derivative of formula I as defined above in claim 1, for treatment of pain.

- 6. (currently amended) A composition method, use or package according to claim 1 any one of the preceding claims in which the COX-2 inhibitor is selected from the group consisting of rofecoxib, etoricoxib, celecoxib, valdecoxib, parecoxib, er and a 5-alkyl-2-arylaminophenylacetic acid derivative COX-2 inhibitor, or a pharmaceutically acceptable salt thereof, or any hydrate thereof.
- (currently amended) A composition method, use or package according to claim 1
 [[7]] in which the COX-2 inhibitor is a compound of formula V

$$\begin{array}{c|c} R & CH_2COOH \\ \hline NH & R_5 \\ \hline R_2 & R_4 \\ \hline R_3 & \\ -3- \end{array}$$

wherein R is methyl or ethyl;

R₁ is chloro or fluoro;

R₂ is hydrogen or fluoro;

R₃ is hydrogen, fluoro, chloro, methyl, ethyl, methoxy, ethoxy or hydroxy;

R₄ is hydrogen or fluoro; and

R₅ is chloro, fluoro, trifluoromethyl or methyl,

or a pharmaceutically acceptable salt or ester thereof.

 (currently amended) A composition method, use or package according to claim 7 in which the COX-2 inhibitor is 5-methyl-2-(2'-chloro-6'-fluoroanilino)phenylacetic acid,

or a pharmaceutically acceptable salt or ester thereof.

9. (currently amended) A composition method, use or package according to claim 1 any one of the preceding claims in which the benzothiadiazole derivative is 5-chloro-4-(2-imidazol-2-ylamino)-2,1,3-benzothiadiazole.